

93206

From: McGarry, Sean
Sent: Monday, May 05, 2003 2:46 PM
To: STIC-Biotech/ChemLib
Subject: SEQ SEARCH

Sean McGarry 73484
AU 1635
CM1 11D07 Office
CM1 11E12 Mailbox
305-7028
09/917,963

Please, a length limited search of nucleotides 3050-3250 of SEQ ID NO:3 (nt≤100).

Thanks

OFF

Searcher: _____
Phone: _____
Location: _____
Date Picked Up: 5/6

Searcher Prep/Review: _____
Clerical: _____
Online Time: _____

TYPE OF SEARCH:

NA Sequences: _____
AA Sequences: _____
Structures: _____
Bibliographic: _____

Full text: _____
Patent Family: _____
Other: _____

VENDOR/COST (where applic.)

STN: _____
DIALOG: _____
Questel/Orbit: _____
DRLink: _____

Sequence Sys.: _____
WWW/Internet: _____
Other (specify): _____



McGarry, Sean

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Please, a length limited search of nucleotides 3050-3250 of SEQ ID NO:3 (nt \leq 100).

Thanks



Urbani, N. I. and A. I. Ponomarev. 1969. *Tr. Gos. univ. Ser. Estest. Nauki* 13: 103-107.

| | | | | |
|-----------------------|-------|----------------|-------|------------|
| 500 pin Match | 14.4% | 500 pin 19 | 15.1% | 100 pin 19 |
| Best Local Similarity | 66.1% | Best No. 10002 | | |



Sequence version 5.1.4-p5-457a
Copyright (c) 1993 - 2003 Computer Tech

Multiple sequence search using SW model

Mon May 12 2003, 00:22:59, Search time 477 seconds

(without alignment)

4431.927 Million cell updates/sec

US-09-917-963-3-copy_3050_3250

Sequence 201

Scoring Matrix

Gapop 10.0, Gapext 1.0

Searched: 16154066 seqs, 8037743376 residues

Total number of hits satisfying chosen parameters: 52784

Minimum DB seq length: 0

Maximum DB seq length: 100

Post processing: Minimum Match 68

Maximum Match 100%

Listing first 45 summaries

FASTA:

1: cm_ost1a.*

2: cm_ost1b.*

3: cm_ost1c.*

4: cm_ost1d.*

5: cm_ost1e.*

6: cm_ost1f.*

7: cm_ost1g.*

8: cm_ost1h.*

9: cm_ost1i.*

10: cm_ost1j.*

11: cm_ost1k.*

12: cm_ost1l.*

13: cm_ost1m.*

14: cm_ost1n.*

15: cm_ost1o.*

16: cm_ost1p.*

17: cm_ost1q.*

18: cm_ost1r.*

19: cm_ost1s.*

20: cm_ost1t.*

Prod. No. is the number of results predicted by Clustal to have a score greater than or equal to the score value indicated, and is derived by analysis of the total score distribution.

SUMMARIES

| Prod. No. | Score | Match | Length | DB | ID | Description |
|-----------|-------|-------|--------|----|-----------|-------------------|
| 1 | 40 | 19.9 | 100 | 1 | 88004554 | AT32451 MR3-BN012 |
| 2 | 42.4 | 18.6 | 74 | 2 | BSR004184 | AT32451 MR3-BN012 |
| 3 | 42.4 | 18.6 | 74 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 4 | 40.4 | 18.1 | 67 | 2 | BSR004184 | AT32451 MR3-BN012 |
| 5 | 40.4 | 18.1 | 69 | 10 | BSR014574 | AT32451 MR3-BN012 |
| 6 | 40.4 | 17.8 | 95 | 9 | BSR014574 | AT32451 MR3-BN012 |

| Prod. No. | Score | Match | Length | DB | ID | Description |
|-----------|-------|-------|--------|----|-----------|-------------------|
| 7 | 45.6 | 17.7 | 88 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 8 | 45.6 | 17.7 | 88 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 9 | 45.6 | 17.7 | 88 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 10 | 45.6 | 17.7 | 88 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 11 | 44.8 | 17.4 | 68 | 9 | BSR014574 | AT32451 MR3-BN012 |
| 12 | 44.8 | 17.4 | 68 | 9 | BSR014574 | AT32451 MR3-BN012 |
| 13 | 44.8 | 17.4 | 68 | 9 | BSR014574 | AT32451 MR3-BN012 |
| 14 | 44.8 | 17.4 | 68 | 9 | BSR014574 | AT32451 MR3-BN012 |
| 15 | 44.8 | 17.4 | 68 | 9 | BSR014574 | AT32451 MR3-BN012 |
| 16 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 17 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 18 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 19 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 20 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 21 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 22 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 23 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 24 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 25 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 26 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 27 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 28 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 29 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 30 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 31 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 32 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 33 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 34 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 35 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 36 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 37 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 38 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 39 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 40 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 41 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 42 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 43 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 44 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 45 | 44.4 | 17.1 | 95 | 1 | BSR014574 | AT32451 MR3-BN012 |

REFERENCES

| Prod. No. | Score | Match | Length | DB | ID | Description |
|-----------|-------|-------|--------|----|-----------|-------------------|
| 1 | 40 | 19.9 | 100 | 1 | 88004554 | AT32451 MR3-BN012 |
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| 3 | 42.4 | 18.6 | 74 | 1 | BSR014574 | AT32451 MR3-BN012 |
| 4 | 40.4 | 18.1 | 67 | 2 | BSR004184 | AT32451 MR3-BN012 |
| 5 | 40.4 | 18.1 | 69 | 10 | BSR014574 | AT32451 MR3-BN012 |
| 6 | 40.4 | 17.8 | 95 | 9 | BSR014574 | AT32451 MR3-BN012 |


```

XX 26 APR 2001: 2001W0 D011419.
XX
XX 27 APR 2000: 2000W0 D001710.
XX
XX (GEO) (GEO) (GEO)
XX
XX Phil A., Tolman R., Glynnov S., Matray L.
XX
XX W01: 2002 02116/03.
XX
XX Use of substituted flavone or isoflavone compounds for inhibiting
XX telomerase activity, and treating e.g. cancer
XX
XX
XX Example 4: Flav-42: English.
XX
XX The present sequence is that of a labelled probe used in a telomerase
XX detection and measurement assay used to test the efficacy of the
XX compounds of the invention as telomerase inhibitors. The specification
XX describes a method of inhibiting telomerase activity in a cell,
XX comprising contacting the cell with a flavone or isoflavone compound. The
XX invention has cytotoxic and ipsosidic, antiproliferative and antitumor
XX immunosuppressive, proto-oncogenic and immunoparasitocidal properties. The compounds
XX of the invention act as telomerase inhibitors and are used for treating
XX telomerase mediated conditions or diseases, e.g. tumours. They may also
XX be used to treat hyperproliferative or autoimmune disorders, such as
XX psoriasis, rheumatoid arthritis, immune system disorders regulating
XX system suppression, immune system reactions to pollen, ivy or pollen cast
XX also comatose, protozoan and fungal infections. The compounds can be
XX administered with other active agents, e.g. anti cancer agents. The
XX compounds can also be administered to plants and soil infected with
XX phyto-pathogenic organisms having telomerase activity, alone or in
XX combination with other agents to control plant diseases. The compounds
XX can be used for a wide variety of malignant cell types and are highly
XX selective, avoiding the problems of current cancer treatments, which are
XX non specific and toxic.
XX
XX Sequence 71 BP: 56 A: 12 C: 9 G: 3 T: 0 other:
XX
XX Query Match 16,98; Score 44; DB 24; Length 71;
XX Post local similarity 74,18; Pred. No. 1,8e-02;
XX Matches 43; Conservative 0; Mismatches 15; Indels 0; Gaps 0
XX
XX 1 to 60AAACACCTTCCTCTCAATACGAAAAAAAAAAAAAAAAAAATACACACACAAATAAA 194
XX 111111111111111111111111111111111111111111111111111111111
XX 2 CTAACCTTAACCTTAACTCAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 59
XX
XX RESULT 2
XX A128139/c
XX
XX A128139 standard; DNA; 83 BP.
XX
XX A128139:
XX
XX 06 JAN 1997 (first entry)
XX
XX Source: gene-related gene sequence 1011, 413.
XX
XX
XX Human fibroblast; AIDS; enhanced differential display; mRNA preparation;
XX sequence of cells; fibroblast cells; defining cell; sequence related to
XX gene expression; non-susceptible cell; age related lipofuscin; retina; EBN;
XX fibroblast; liver spot; donor tissue; suscept melanocyte; melanin;
XX hypomelanation; ss.
XX
XX Synthesizer
XX
XX W0013010 A:
XX
XX 09 MAY 1997:
XX
XX 24 MAY 1997: 96W0-1811230.
XX
XX 31 OCT 1994: 9408-038420.
XX
XX

```

(CHRO)-TEGEN CORP.

F1 Part 1, Park W., Hirsch KS., Linkens MK., Villeportou B.
F1 West MD
XX
XX WP: 1997-051464/25,
FE
X1 Identifying, isolating and regulating sequence-related genes -
F1 used to ameliorate problems associated with accumulation of
F1 senescent cells, e.g. age-related lipofuscin accumulation in the
F1 retina and Ailes

XX
XX Claim 8; Page 46; 135pp; English.

XX
XX AA128076-128113, and AA128141-128173 represent novel senescent-related
CC gene sequences isolated from fibroblasts using the method of the
CC invention. In the method of the invention, mRNA is isolated from a
CC senescent cell, and a young quiescent cell, and the mRNAs are amplified
CC (using primers such as those shown in AA128044-128072) in separate
CC reaction mixtures. The amplified sequences are then separated by size or
CC charge, and the products are analysed to identify a gene from young
CC quiescent cells and dividing cells, that is present at a different level
CC from senescent cells. To enhance the method even more, it can be
CC performed in conjunction with an enhanced differential display (EDD)
CC method (an mRNA preparation method) on the fibroblasts. The method can
CC be used for the rapid and efficient identification and isolation of
CC senescent-related genes and gene products, and to detect and distinguish
CC between senescent and non-senescent cells. It can also be used to
CC identify cells expressing senescence specific (or related) gene products,
CC and to screen for compounds capable of altering gene expression in
CC senescent cells. The method can also be used to ameliorate problems
CC associated with the accumulation of senescent cells such as age-related
CC fibrosis with accumulation in the retina, and in the treatment of AIDS.
CC Also, the method can be used to distinguish young cells from senescent
CC cells in donor tissue, which is useful in removing senescent melanocytes
CC overexpressing melanin which cause hyper-pigmentation, or liver spots.

XX
XX Sequence 84 BP; 20 A; 5 G; 10 C; 48 T; 0 other;

SO Query Match 16,98; Score 34; DB 17; Length 83;
Percent Similarity 53.4%; Pred. No. 1,807,02;
Matches 52; Conservative 0; Mismatch 30; Indels 0; Gaps 0

CY 100 GTCTGATAGCAATTGTCTTAACATGACGACAAAACAAACGTTTCTTAATTCACCAA 159
1 111 1111 11111111 11 1111111111 111 11111111
DB 82 GTCTCATATACACTTCTTGTAACAAGCAATTAATAAATTAAAGTATTCTTAAAAAAA 24
1 111111111111111111 11
CY 160 AAAAAAAAAAAAAAAAAAAATAA 161
111111111111111111 11
DB 22 AAAAAAAAAAAAAAAAAAAAAA 1

RESUIT 4
AAC13249
ID AAC13249 standard; cDNA; 81 BP.
XX
XX AAC13249;
IT 36 OCT 2000 (first entry)
XX
XX Human secreted protein c' PST, SPQ ID NO. 17324.
FE
XX Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;
XX gene therapy; chromosome mapping; SS:
OS Homo sapiens.
XX
XX EP034401-A2.
IN
XX
XX 06-SEP-2000.
XX
XX 21 MAR 2000; 200001-020040.

Author: Thomas E. Welton

Title:

BL44: 4, Room 120, NIH Campus, Bethesda, MD 20892-0425

Phone: (301) 496-4521

Fax: (301) 496-4521

Email: tomwel@nih.gov

Project: A: A7AATGTAATAATG

Project: B: ATCTTCTTTTATG

Seq. size: 90

Seq. type: 90

Seq. type: 90

Seq. type: 90

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Seq. type: 90

Seq. type: 90

Buffer: 10 mM Tris-HCl, pH 8.3

50 mM KCl

50 mM MgCl₂

1 mM DTT

1 mM EDTA

1 mM EGTA

1 mM MgSO₄

1 mM CaCl₂

1 mM MnCl₂

1 mM ZnCl₂

1 mM CuCl₂

1 mM NiCl₂

1 mM CoCl₂

1 mM FeCl₃

1 mM AlCl₃

1 mM GaCl₃

1 mM InCl₃

1 mM SnCl₄

1 mM SbCl₅

1 mM TeCl₆

1 mM BiCl₃

1 mM PbCl₂

1 mM HgCl₂

1 mM AsCl₃

1 mM SeCl₆

1 mM BrCl₃

1 mM ICl₃

1 mM AtCl₃

1 mM TlCl₃

1 mM PbCl₂

1 mM BiCl₃

1 mM SbCl₅

1 mM TeCl₆

1 mM BrCl₃

1 mM ICl₃

1 mM AtCl₃

1 mM TlCl₃

1 mM PbCl₂

1 mM BiCl₃

1 mM SbCl₅

1 mM TeCl₆

1 mM BrCl₃

10 mM Tris-HCl, pH 8.3

50 mM KCl

50 mM MgCl₂

1 mM DTT

1 mM EDTA

1 mM EGTA

1 mM MgSO₄

1 mM CaCl₂

1 mM MnCl₂

1 mM ZnCl₂

1 mM CuCl₂

1 mM NiCl₂

1 mM CoCl₂

1 mM FeCl₃

1 mM AlCl₃

1 mM GaCl₃

1 mM InCl₃

1 mM SnCl₄

1 mM SbCl₅

1 mM TeCl₆

1 mM BrCl₃

1 mM ICl₃

1 mM AtCl₃

1 mM TlCl₃

1 mM PbCl₂

1 mM BiCl₃

1 mM SbCl₅

1 mM TeCl₆

1 mM BrCl₃

1 mM ICl₃

1 mM AtCl₃

1 mM TlCl₃

1 mM PbCl₂

1 mM BiCl₃

1 mM SbCl₅

1 mM TeCl₆

1 mM BrCl₃

1 mM ICl₃

1 mM AtCl₃

1 mM TlCl₃

1 mM PbCl₂

1 mM BiCl₃

10 mM Tris-HCl, pH 8.3

50 mM KCl

50 mM MgCl₂

1 mM DTT

1 mM EDTA

1 mM EGTA

1 mM MgSO₄

1 mM CaCl₂

1 mM MnCl₂

1 mM ZnCl₂

1 mM CuCl₂

1 mM NiCl₂

1 mM CoCl₂

1 mM FeCl₃

1 mM AlCl₃

1 mM GaCl₃

1 mM InCl₃

1 mM SnCl₄

1 mM SbCl₅

1 mM TeCl₆

1 mM BrCl₃

1 mM ICl₃

1 mM AtCl₃

1 mM TlCl₃

1 mM PbCl₂

1 mM BiCl₃

1 mM SbCl₅

1 mM TeCl₆

1 mM BrCl₃

1 mM ICl₃

1 mM AtCl₃

1 mM TlCl₃

1 mM PbCl₂

1 mM BiCl₃

1 mM SbCl₅

1 mM TeCl₆

1 mM BrCl₃

1 mM ICl₃

1 mM AtCl₃

1 mM TlCl₃

1 mM PbCl₂

1 mM BiCl₃

10 mM Tris-HCl, pH 8.3

50 mM KCl

50 mM MgCl₂

1 mM DTT

1 mM EDTA

1 mM EGTA

1 mM MgSO₄

1 mM CaCl₂

1 mM MnCl₂

1 mM ZnCl₂

1 mM CuCl₂

1 mM NiCl₂

1 mM CoCl₂

1 mM FeCl₃

1 mM AlCl₃

1 mM GaCl₃

1 mM InCl₃

1 mM SnCl₄

1 mM SbCl₅

1 mM TeCl₆

1 mM BrCl₃

1 mM ICl₃

1 mM AtCl₃

1 mM TlCl₃

1 mM PbCl₂

1 mM BiCl₃

1 mM SbCl₅

1 mM TeCl₆

1 mM BrCl₃

1 mM ICl₃

1 mM AtCl₃

1 mM TlCl₃

1 mM PbCl₂

1 mM BiCl₃

1 mM SbCl₅

1 mM TeCl₆

1 mM BrCl₃

1 mM ICl₃

1 mM AtCl₃

1 mM TlCl₃

1 mM PbCl₂

1 mM BiCl₃

10 mM Tris-HCl, pH 8.3

50 mM KCl

50 mM MgCl₂

1 mM DTT

1 mM EDTA

1 mM EGTA

1 mM MgSO₄

1 mM CaCl₂

1 mM MnCl₂

1 mM ZnCl₂

1 mM CuCl₂

1 mM NiCl₂

1 mM CoCl₂

1 mM FeCl₃

1 mM AlCl₃

1 mM GaCl₃

1 mM InCl₃

1 mM SnCl₄

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1 mM AtCl₃

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1 mM BiCl₃

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1 mM TeCl₆

1 mM BrCl₃

1 mM ICl₃

1 mM AtCl₃

1 mM TlCl₃

